AURICAL FreeFit

AURICAL FreeFit and the OTOsuite PMM Module

User Guide

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Technical support
Please contact your supplier.

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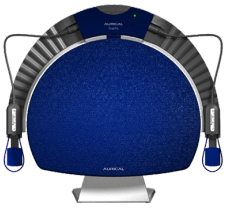
1 Device Description

FreeFit is used for measuring the sound pressure level close to the client's eardrum as well as outside the ear by the pinna for both ears simultaneously.

AURICAL FreeFit should only be charged using the type 1053 charger unit or the type 1081 AURICAL speaker unit from GN Otometrics A/S.

Type 1053 charger unit



AURICAL speaker unit with charger

Note • *This manual describes the assembly and use of the type 1053 charger unit. If you use the AURICAL speaker unit to charge your FreeFit, see the manual for AURICAL Aud.*

Note • *For information about the Counseling and Simulations software, see the manual for AURICAL Visible Speech and the Counseling and Simulations Module.*

You use the OTOsuite PMM module and the Counseling and Simulations module to operate FreeFit. FreeFit cannot be used without OTOsuite software.

2 Intended use

Users: audiologists, hearing instrument dispensers, ENTs, speech therapists and other health care professionals.

Use: to visualize the amplified signal recorded in the ear with reference information such as target curves to provide an objective basis for adjusting the hearing instrument settings.

2.1 Typographical conventions

The use of Warning, Caution and Note

For safety reasons and appropriate use of the device and/or software, the manual contains **Warnings**, **Cautions** and **Notes**. These headings are used as follows:

Warning • *Indicates that there is risk of death or serious injury to the user or patient.*

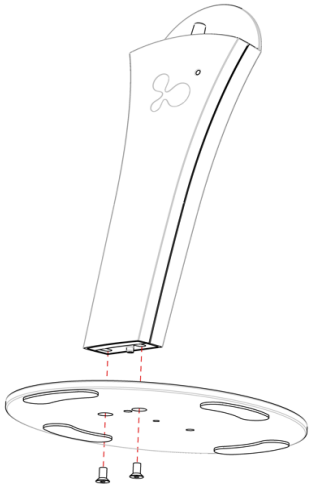
Caution • *Indicates that there is a risk of injury to the user or patient or risk of damage to data or the device.*

Note • *Indicates that you should take special notice.*

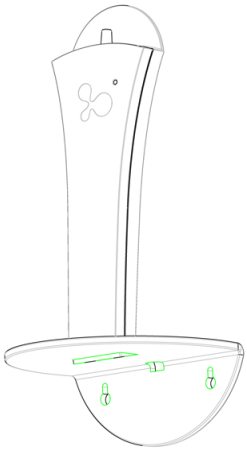
3 Unpacking FreeFit

1. Unpack the device carefully.
   When you unpack the device and accessories, it is a good idea to keep the packing material in which they were delivered. If you need to send the device in for service, the original packing material will protect against damage during transport, etc.
2. Visually inspect the equipment for possible damage.
   If damage has occurred, do not put the device into operation. Contact your local distributor for assistance.
3. Check with the packing list to make sure that you have received all necessary parts and accessories. If your package is incomplete, contact your local distributor.

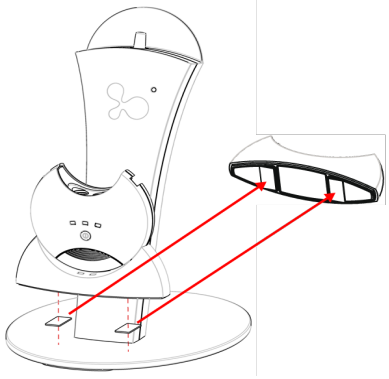
4 Installation



Installation for desk top use



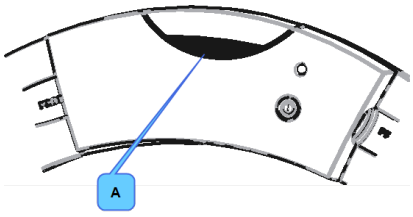
Wall mount installation



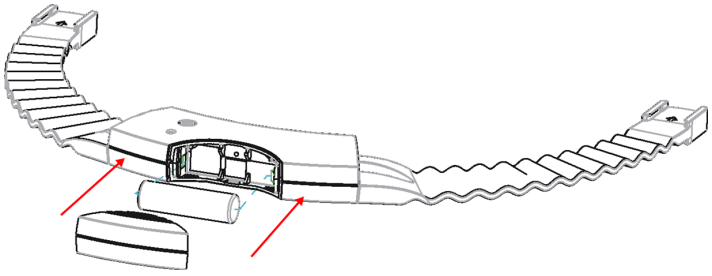
Mounting the NOAHLink charger on the FreeFit charger base plate

5 Powering the device

Caution • *Use only the following battery types:*
Rechargeable (Ni-MH type) AA (R6) 1.2V, 1 pc. (Use only rechargeable batteries supplied by GN Otometrics A/S)
Alkaline AA (R6) 1.2V, 1 pc.



A. Press to open



5.1 Recharging the battery using the charger stand

Warning • *If you are using an alkaline battery, do not attempt to charge your AURICAL FreeFit. Your alkaline battery may be damaged and leak, and this may in turn cause damage to FreeFit. Place FreeFit in the charger unit only if FreeFit contains a rechargeable battery.*
Batteries should be removed if equipment is not likely to be used for some time.

Caution • *To power the charger, you must use an IEC/UL 60601-01 certified power adaptor supplying 9 V DC, min. 300 mA and with a maximum available output of 15 W. The adapter supplied with the unit meets these specifications.*

6 Switching FreeFit on or off

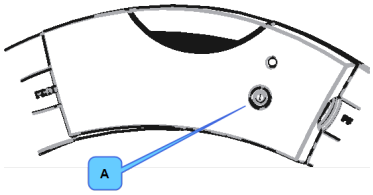
> ***Warning*** • *Unless you are charging FreeFit with the AURICAL Aud speaker unit, which has a medically isolated power supply unit, do not attempt to use AURICAL FreeFit with clients while it is placed in the charger unit.*

To switch on FreeFit

Press and hold the power button on top of the device until the status indicator light turns on. The status indicator will light for about 3 seconds, and then go into periodic flashing.

To switch off FreeFit

Press and hold the power button on top of the device until the status indicator light turns off.



A. Power button

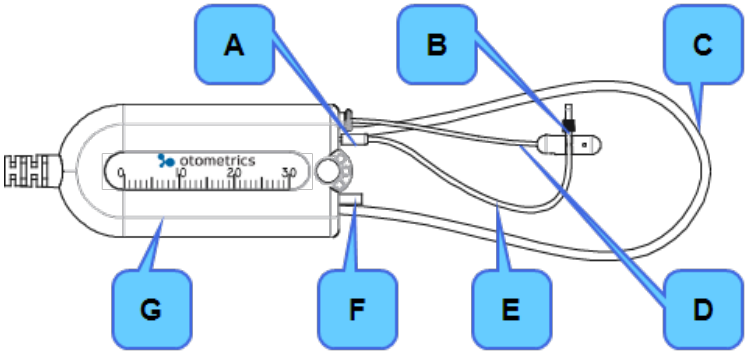
7 Connecting FreeFit to PMM

When you use PMM for the first time, run the configuration wizard to set up the connection between FreeFit and PMM.

After you have configured PMM for the first time, if FreeFit is turned on when you open the Control Panel in PMM, then FreeFit will connect to PMM automatically. Otherwise, you can connect FreeFit as follows:

1. Switch on FreeFit.
2. In PMM, on the toolbar, click **Control Panel**.
3. In the control panel, click **Connect**.

8 FreeFit probes



A. Probe tube port
B. Marker ring
C. Ear cord
D. Tube guide
E. Probe tube
F. Transducer tube port (RECD probe only)
G. Probe housing

The ear cord is used to hang the probe below the client's ear. The probe tube is inserted into the ear canal for probe microphone measurements. The probe tube has a black marker ring for marking how far into the ear canal the tube should be inserted. The tube guide is used to stabilize the position of the probe tube. Before you make RECD measurements, you fit a transducer tube on the transducer tube port.

8.1 Fitting probe tubes on the probes

A bag of silicone probe tubes is supplied together with FreeFit.

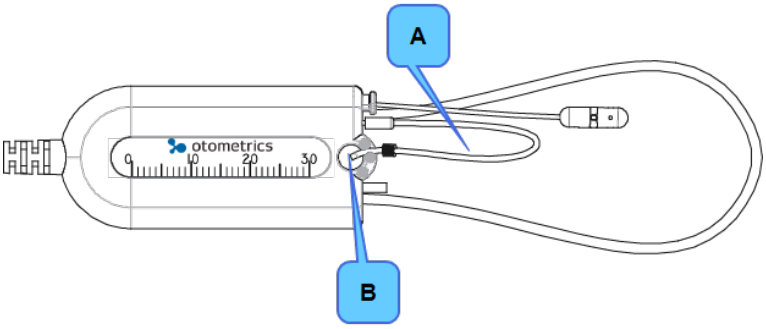
To fit a probe tube on the probe

Fit a probe tube to the probe tube port (thin metal tube) at the top of the probe housing. Gently push and twist the probe tube down as far as possible over the port.

8.2 Calibrating the probe tubes

> ***Note*** • *To prevent cross-infection, use new probe tubes for each client.*

1. Fit a new probe tube on the probe.
2. Insert the free end of the probe tube in the test location on the probe.



A. Probe tube
B. Test location for probe tube calibration

3. Make sure that FreeFit is connected to PMM.
4. Press the power button briefly on FreeFit.

   The **Probe Tube Calibration** dialog box appears and the calibration starts automatically.

   Alternatively, launch the wizard with the toolbar icon. (In RECD, pressing the power button starts an ear measurement.)

5. If the tube calibration fails, check whether the tubes are blocked (pinched or clogged) and try to eliminate sources of ambient noise.

8.3 Fitting the probes on the client and inserting the probe tubes

It is important that the probe tube for every measurement is inserted correctly and consistently in the ear of the client.

1. Place the black marker ring at the recommended distance from the tip of the probe tube.

   Recommended distances are

men:	27 mm
women:	27 mm
children:	20-25 mm

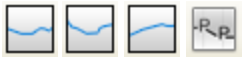
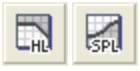
   In the case of children, otoscopy is especially recommended to prevent contact with the eardrum.

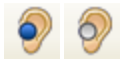
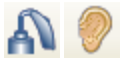
2. Place the ear cords with the probes over the ears of the client. Adjust the length of the ear cords, if required.

3. **Be careful!**
   Carefully insert the probe tube into the ear canal until the black marker ring reaches the intertragal notch.

9 Toolbar icons in PMM

The icons available in the toolbar depend on the test function that you have selected.

Toolbar icons in PMM	
	Select device.
	Toggle between Response, Gain, REIG and Predicted Aided Audiogram (PAA) views.
	Select view: **HL**: Hearing Level or **SPL**: Sound Pressure Level
	Select audiogram view: Left, both or right
	Open **Fitting Details** dialog. ***Note*** • *You must select the **Use OpenREM calibration** option if you are fitting an open ear instrument.*
	Open **Listen at the Eardrum/Listen in the Coupler** window to record the signal at the eardrum or in the coupler, or monitor the signal through your headphones.
	Show/hide **Curve Legend** and **Overlays** box.

Toolbar icons in PMM	
	Switch to **Feature-2-Benefit** view. (FreeStyle only)
	Launch probe tube calibration wizard.
	Toggle between standard calibration and OpenREM calibration.
	Select previously measured RECD values.
	Toggle between coupler fitting mode and Real Ear fitting mode.
	Show/Hide **OnTarget** view, which displays the difference between the target curve and the measured curve.
	Switch to **On Top** mode.

10 Performing probe microphone measurements

When you start a new session in PMM, you must ensure the fitting parameters are set correctly in the **Fitting Details** dialog box.

To set the fitting parameters

1. Press F10 to open the **Fitting Details** dialog box.
2. Select the appropriate target rule.
3. Fill in the remaining fields in the dialog box.

Making Probe Microphone Measurements

The following sections describe the main procedures involved in PMM:

Measuring RECD ► 11
Measuring Unaided Response ► 13
Measuring Occluded Response ► 14
Measuring Aided Response ► 14

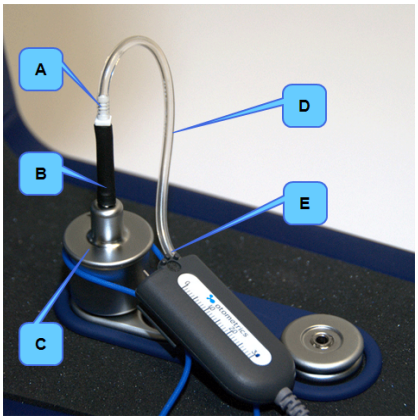
10.1 Measuring RECD

If you want to use measured RECD values for coupler based fitting, you can measure RECD in PMM as follows:

Measure coupler response:

Skip this procedure if you have a stored coupler measurement.

1. Open the **RECD** tab in PMM.
2. Indicate the type of coupler adapter you are using, and whether you are using an ear mold or foam insert tip.
3. Click **Coupler Response...** in the RECD Control Panel.
4. Attach the right RECD ear probe to the coupler in AURICAL HIT.

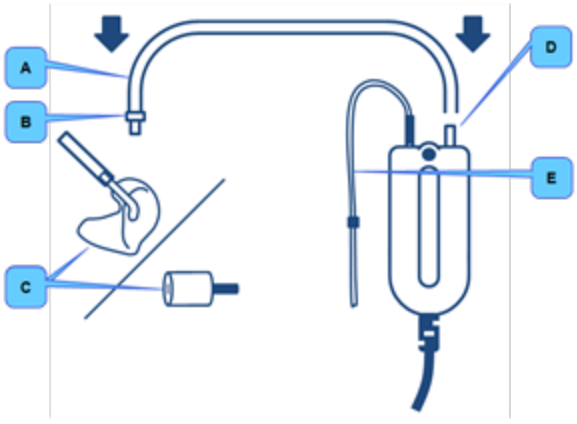


A. RECD coupling
B. BTE adapter tube
C. BTE (HA2) adapter
D. Transducer tubing
E. Transducer tube port

5. Click the **Measure Right** button.
6. Connect the left probe to the coupler in AURICAL HIT.
7. Click the **Measure Left** button.
8. Click **OK**.
9. Remove the probe from AURICAL HIT and remove the RECD coupling from the tubing of the BTE coupler.

Then measure real ear response:

1. Attach the probes to the FreeFit.
2. Perform probe tube calibration.
3. Connect the RECD coupling to the ear mold tubing (or foam insert tip).



A. Transducer tubing
B. RECD coupling
C. Ear mold or foam insert tip
D. Transducer tube port
E. Probe tube

4. Place the probe tubes in the client's ears together with the ear molds or foam insert tips (see Fitting the probes on the client and inserting the probe tubes ▶ 9).

5. Select ear to measure.

6. In the control panel, click **Ear Response** (or briefly press the power button on the FreeFit).

The measured ear response and the RECD are displayed in their respective graphs.

10.2 Measuring Unaided Response

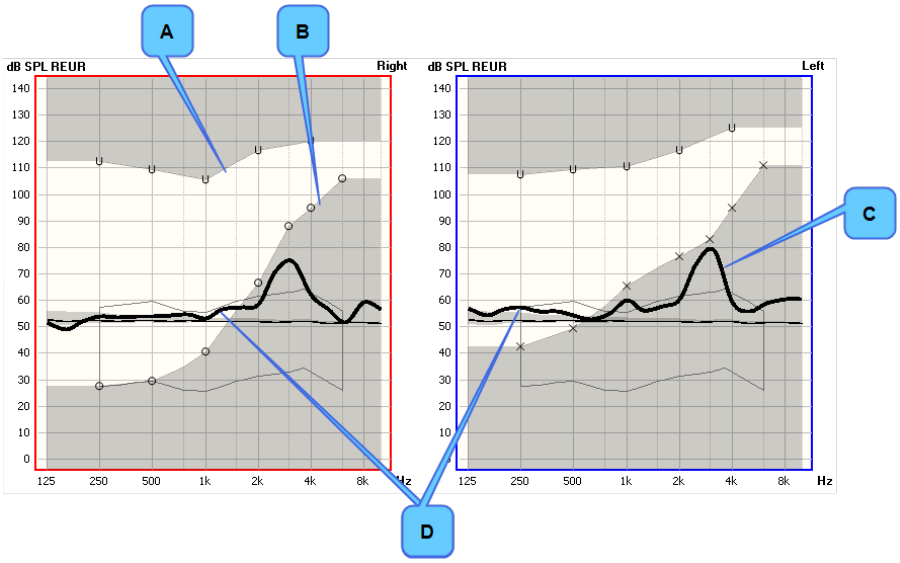
In the **Unaided** screen, measure without hearing instruments to determine the natural amplification of the ear canal.

1. Insert probe tube.
2. Select ear to measure.

3. Select graph.

4. Click the **Unaided** button on the Control Panel.

> ***Note*** • *In an unaided measurement, we usually expect a peak on the measurement curve around the 3kHz frequency of about 10-20 dB SPL.*



A. UCL
B. Audiogram
C. Peak around 3 kHz
D. Measurement curve

10.3 Measuring Occluded Response

In the **Occluded Response** screen, measure with muted hearing instruments in ears to measure the occlusion or openness of the fitting.

1. Place the hearing instrument on the ear of the patient, with the probe tube inserted in the ear canal. Ensure the hearing instrument is muted or turned off.
2. Click the **Occluded** button on the Control Panel.

When you compare the REUR to the REOR, you can see the impact of the occlusion of the ear canal.

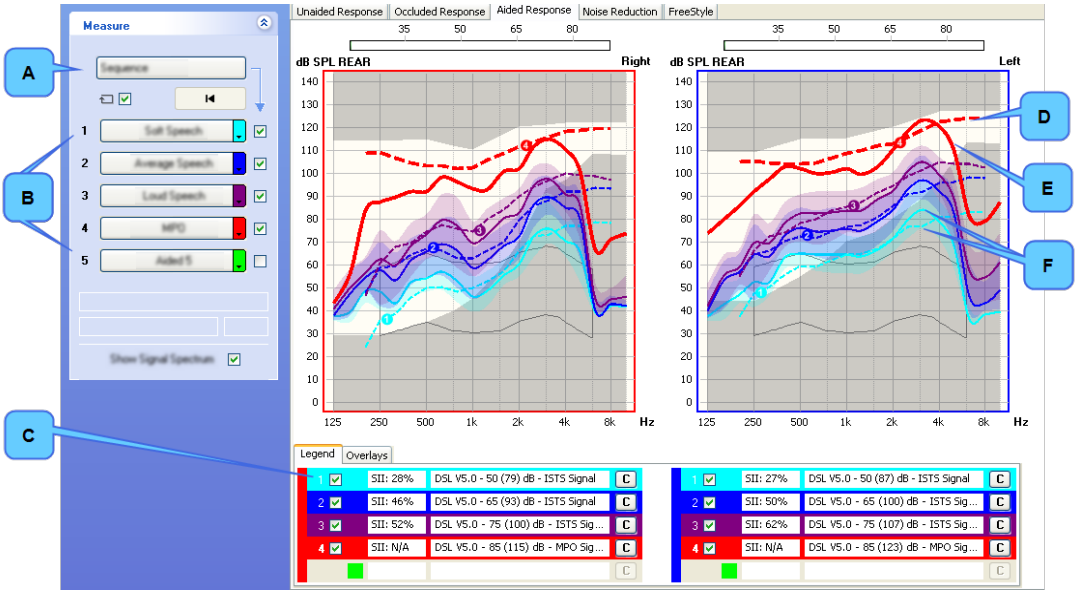
10.4 Measuring Aided Response

In the **Aided** screen, measure the gain that the hearing instruments are providing in relation to a specified prescriptive fitting target.

1. Place the hearing instrument on the ear of the patient, with the probe tube inserted in the ear canal.

> ***Note*** • *For Coupler based fitting, select the Coupler based fitting icon on the toolbar and attach the hearing instrument to the coupler in the AURICAL HIT (see the manual for AURICAL HIT).*

2. Switch the hearing aid on without moving it.
   All the hearing instrument features should be left on with the general use program selected.
3. Configure the control panel to play up to 5 signals. For example: 3 input levels for a speech or speech-like stimulus: Soft (50/55 dB SPL), Average (65 dB SPL) and Loud (80 dB SPL) plus an MPO stimulus.
4. Present the various input levels and MPO signal separately or in one sequence.
5. Compare the measured aided response curve to the prescriptive target values (dashed curve) and the measured MPO curve to the UCL. Consider adjusting the MPO (Maximum Power Output) if indicated by any report of loudness discomfort.
6. Adjust the hearing instrument with the fitting software to achieve the desired gain, and repeat the measurements to evaluate the effects of the changes.



A. Sequence button
B. Measurement buttons for different stimuli
C. Curve legend for button 1.
D. Target curve for MPO.
E. Measurement curve for MPO.
F. Measurement curve and target curve for button 1.

11 Demonstrating hearing instrument features

Use the **Noise Reduction** screen to evaluate and demonstrate the hearing instrument's noise reduction feature. Each Noise Reduction test is an automatic sequence that contains two curves (with a delay between the two curves):

- Curve 1 – a snapshot taken immediately before Noise Reduction takes effect.
- Curve 2 – a snapshot taken automatically after the selected Adaptation Interval, when the Noise Reduction has taken effect.

To demonstrate the Noise Reduction feature:

1. Program the hearing instrument for the desired Noise Reduction setting.

2. Configure the measurement buttons to demonstrate the conditions you prefer. For each button, select the time difference between the two measurements (Curve 1 and Curve 2).
3. Click a measurement button in the control panel.

The snapshot curves are displayed in the graph and the overall Noise Reduction is displayed in the curve legend.

The **Feature-2-Benefit** view gives you the opportunity to see and show the gain difference in an easy to understand graph.

The **FreeStyle** test screen is similar to the other PMM test screens but with numerous possibilities to customize protocols.

12 Service, cleaning and maintenance

> ***Warning*** • *Under no circumstances disassemble FreeFit or the FreeFit charger. Contact your supplier. Parts inside FreeFit and the FreeFit charger must only be checked or serviced by authorized personnel.*

12.1 Service

For the sake of safety and in order not to void the warranty, service and repair of electro-medical equipment should be carried out only by the equipment manufacturer or by service personnel at authorized workshops. In case of any defects, make a detailed description of the defect(s) and contact your supplier. Do not use a defective device.

12.2 Cleaning

Use a soft, slightly damp cloth with a small amount of mild detergent or approved non-caustic medical grade disinfectant wipes to clean the unit and charger according to local infection control regulations.

> ***Caution*** • *Keep the unit away from liquids. Do not allow moisture inside the unit.*

> ***Caution*** • *Never immerse the FreeFit probes into water or other cleaning solutions.*

> ***Caution*** • *No part of FreeFit or its accessories is suitable for autoclaving or thermal disinfection/sterilization methods.*

Probe tubes, tube guides and ear cords

These parts are in constant contact with your clients.

- Probe tubes:
  The only part which is inserted into the ear canal during PMM testing is the Probe tube. These tubes are disposable, and should only be used once per client.
- Tube guides and ear cords:
  Use a soft, slightly damp cloth with a small amount of detergent to clean the cords and the tube guides.

Disposal

There are no special requirements regarding the disposal of the silicone test tubes, i.e. they can be discarded according to local regulations.

12.3 Maintenance

Annual calibration

FreeFit and the FreeFit probes must be calibrated once a year by your authorized service department.

13 Other references

For more information, see the online Help in OTOsuite, which contains detailed reference information about FreeFit and PMM, and other OTOsuite products.

For instructions on installing OTOsuite, see the OTOsuite Installation Guide, which you can find on the OTOsuite installation medium (disk or memory stick).

14 Standards

Safety:	IEC 60601-1, UL 60601-1, CAN/CSA -C22.2 NO 601.1-90 AURICAL FreeFit: IEC 60601-1, Internally Powered, Type BF, IPX0
EMC:	IEC 60601-1-2, EN 300 328-2, EN 301 489-17

15 Power adaptor

BRIDGEPOWER CORP MENB1010A0903B01	Output: 9 VDC, 1.10 A Input: 100-240 V, 50-60 Hz
DONGGUAN SHILONG FUHUA ELECTRONIC CO., LTD. UE08WCP-090056SPA	Output: 9 VDC, 0.56 A Input: 100-240 V, 50-60 Hz

Use only the power adaptor supplied with the instrument.

16 Operating environment

Temperature:	+15°C to +35°C (59°F to +95°F)
Relative humidity:	30 to 90%, non-condensing
Warm-up time:	< 1 min.
Air pressure:	600 hPa to 1060 hPa

Operation at temperatures below -20°C or above +60°C may cause permanent damage.

17 Storage and handling

Temperature:	-20°C to +60°C (-4°F to +140°F)
Rel. humidity:	< 90%, non-condensing
Air pressure:	500 hPa to 1060 hPa

18 Manufacturer

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18.1 Responsibility of the manufacturer

The manufacturer is to be considered responsible for effects on safety, reliability, and performance of the equipment only if:

- All assembly operations, extensions, re-adjustments, modifications or repairs are carried out by the equipment manufacturer or personnel authorized by the manufacturer.
- The electrical installation to which the equipment is connected complies with EN/IEC requirements.
- The equipment is used in accordance with the instructions for use.

The manufacturer reserves the right to disclaim all responsibility for the operating safety, reliability and performance of equipment serviced or repaired by other parties.

19 Warning notes

This manual contains information and warnings, which must be followed to ensure the safe performance of the devices and software covered by this manual. Local government rules and regulations, if applicable, should also be followed at all times.

AURICAL FreeFit should only be charged using the type 1053 charger unit or the type 1081 AURICAL speaker unit from GN Otometrics A/S.

Unless you are charging FreeFit with the AURICAL Aud speaker unit, which has a medically isolated power supply unit, do not attempt to use AURICAL FreeFit with clients while it is placed in the charger unit.

1. There are no user-serviceable parts inside the cabinet of the device or charger. For the sake of safety and in order not to void the warranty, service and repair of electro-medical equipment should be carried out only by the equipment manufacturer or by service personnel at authorized workshops. In case of any defects, make a detailed description of the defect(s) and contact your supplier. Do not use a defective device.
2. Keep the unit away from liquids. Do not allow moisture inside the unit.
3. Do not use the instrument in the presence of flammable anesthetics (gases).
4. Unwanted noise may occur if the instrument is exposed to a strong radio field. Such noise may interfere with the process of recording an accurate audiogram and fitting a hearing instrument. Many types of electrical devices, e.g. mobile telephones, may generate radio fields. We recommend that the use of such devices in the vicinity of this instrument be restricted as much as possible.
5. Changes or modifications not expressly approved by the manufacturer could void the user's authority to operate the equipment.
6. This equipment has been tested and found to comply with the limits for a Class B digital device, pursuant to part 15 of the FCC Rules. These limits are designed to provide reasonable protection against harmful interference in a residential installation. This equipment generates, uses and can radiate radio frequency energy and, if not installed and used in accordance with the instructions, may cause harmful interference to radio communications. However, there is no guarantee that interference will not occur in a particular installation. If this equipment does cause harmful interference to radio or television reception, which can be determined by turning the equipment off and on, the user is encouraged to try to correct the interference by one or more of the following measures:
   - Reorient or relocate the receiving antenna.
   - Increase the separation between the equipment and receiver.
   - Connect the equipment into an outlet on a circuit different from that to which the receiver is connected.
   - Consult the dealer or an experienced radio/TV technician for help.
7. For use in Canada: To prevent radio interference to the licensed service, this device is intended to be operated indoors and away from windows to provide maximum shielding. Equipment (or its transmit antenna) that is installed outdoors is subject to licensing.
8. No parts may be eaten, burnt, or in any way used for purposes other than the applications defined in the Intended Use section of this manual.
9. The device and charger unit can be disposed of as normal electronic waste, according to local regulations. Please investigate local regulations concerning the disposal of rechargeable and alkaline batteries.
10. For safety reasons, accessories connected to the equipment's outlet fittings must be identical to the type supplied with the system.

11. It is recommended that an annual calibration be performed on accessories containing transducers. Furthermore, it is recommended that calibration be performed if the equipment has suffered any potential damage (e.g. headphones dropped on the floor).
12. To comply with EN 60601-1-1 computer and printer must be placed out of reach of the client, i.e. not closer than approx. 1.5 meters/5 ft.
13. In the United States of America, Federal law restricts this device to sale by or on the order of a licensed physician.
14. It is recommended to install the unit in an environment that minimizes the amount of static electricity. For example, anti-static carpeting is recommended.
15. The charger unit should be kept away from the client area.

20 Notes on EMC (Electromagnetic Compatibility)

- AURICAL FreeFit is part of a medical electrical system and is thus subject to special safety precautions. For this reason, the installation and operating instructions provided in this document should be followed closely.
- Portable and mobile high-frequency communication devices, such as mobile phones, may interfere with the functioning of AURICAL FreeFit.

<table>
<tr><td colspan="3">Guidance and manufacturer's declaration - electromagnetic emissions for all equipment and systems</td></tr>
<tr><td colspan="3">AURICAL FreeFit is intended for use in the electromagnetic environment specified below. The user of AURICAL FreeFit should ensure that it is used in such an environment.</td></tr>
<tr><td>Emissions test</td><td>Compliance</td><td>Electromagnetic environment - guidance</td></tr>
<tr><td>RF emissions
CISPR 11</td><td>Group 1</td><td>AURICAL FreeFit uses RF energy only for its internal function. Therefore, its RF emissions are very low and are not likely to cause any interference in nearby electronic equipment.</td></tr>
<tr><td>RF emissions
CISPR 11</td><td>Class B</td><td rowspan="3">AURICAL FreeFit is suitable for use in all environments, including domestic environments and those directly connected to the public low-voltage power supply network that supplies buildings used for domestic purposes.</td></tr>
<tr><td>Harmonic emissions IEC 61000-3-2</td><td>Not applicable</td></tr>
<tr><td>Voltage fluctuations/flicker emissions IEC 61000-3-3</td><td>Not applicable</td></tr>
</table>

Guidance and manufacturer's declaration - electromagnetic immunity for all equipment and systems			
AURICAL FreeFit is intended for use in the electromagnetic environment specified below. The user of AURICAL FreeFit should ensure that it is used in such an environment.			
Immunity test	**IEC 60601 test level**	**Compliance level**	**Electromagnetic environment - guidance**
Electrostatic discharge (ESD) IEC 61000-4-2	+/- 6 kV contact +/- 8 kV air	+/- 6 kV contact +/- 8 kV air	Floors should be wood, concrete or ceramic tile. If floors are covered with synthetic material, the relative humidity should be at least 30%.
Power frequency (50/60 Hz) magnetic field IEC 61000-4-8	3 A/m	3 A/m	Power frequency magnetic fields should be at levels characteristic of a typical location in a typical commercial or hospital environment.
Note • U_T is the AC mains voltage prior to application of the test level.			

Guidance and manufacturer's declaration - electromagnetic immunity - for equipment and systems that are NOT life-supporting			
AURICAL FreeFit is intended for use in the electromagnetic environment specified below. The user of AURICAL FreeFit should ensure that it is used in such an environment.			
Immunity test	IEC 60601 test level	Compliance level	Electromagnetic environment - guidance
Radiated RF IEC 61000-4-3	3 V/m 80 MHz to 2.5 GHz	3 V/m	Portable and mobile RF communications equipment should be used no closer to any part of AURICAL FreeFit, including cables, than the recommended separation distance calculated from the equation applicable to the frequency of the transmitter. Recommended separation distance: $d = 1.2\sqrt{P}$ for 80 MHz to 800 MHz $d = 2.3\sqrt{P}$ for 80 MHz to 2.5 GHz, where *P* is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer and d is the recommended separation distance in metres (m). Field strengths from fixed RF transmitters, as determined by an electromagnetic site survey, [a] should be less than the compliance level in each frequency range. [b] Interference may occur in the vicinity of equipment marked with this symbol:
Note 1: At 80 MHz and 800 MHz the separation distance for the higher frequency range applies. **Note 2**: These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects and people.			
a. Field strengths from fixed transmitters, such as base stations for radio (cellular/cordless) telephones and land mobile radios, amateur radio, AM and FM radio broadcast and TV broadcast cannot be predicted theoretically with accuracy. To assess the electromagnetic environment due to fixed RF transmitters, an electromagnetic site survey should be considered. If the measured field strength in the location in which AURICAL FreeFit is used exceeds the applicable RF compliance level above, the AURICAL FreeFit should be observed to verify normal operation. If abnormal performance is observed, additional measures might be necessary, such as reorienting or relocating AURICAL FreeFit. b. Over the frequency range 150 kHz to 80 MHz, field strengths should be less than 3 V/m.			

Recommended separation distances between portable and mobile RF communications equipment and AURICAL FreeFit		
The AURICAL FreeFitis intended for use in an electromagnetic environment in which radiated RF disturbances are controlled. The customer or the user of the AURICAL FreeFit can help prevent electromagnetic interference by maintaining a minimum distance between portable and mobile RF communications equipment (transmitters) and the AURICAL FreeFitas recommended below, according to the maximum output power of the communications equipment.		
Rated maximum output power of transmitter W	Separation distance according to frequency of transmitter m	
	80 MHz to 800 MHz $d = 1.2\sqrt{P}$	800 MHz to 2,5 GHz $d = 2.3\sqrt{P}$
0.01	0.12	0.23
0.1	0.38	0.73
1	1.2	2.3
10	3.8	7.3
100	12	23
For transmitters rated at a maximum output power not listed above, the recommended separation distance d in metres (m) can be estimated using the equation applicable to the frequency of the transmitter, where *P* is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer. **Note 1**: At 80 MHz and 800 MHz the separation distance for the higher frequency range applies. **Note 2**: These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects and people.		

21 Definition of symbols

	Electronic equipment covered by the Directive 2002/96/EC on waste electrical and electronic equipment (WEEE). All electrical and electronic products, batteries, and accumulators must be taken to separate collection at the end of their working life. This requirement applies in the European Union. Do not dispose of these products as unsorted municipal waste. You can return your device and accessories to Otometrics, or to any Otometrics supplier. You can also contact your local authorities for advice on disposal.
	Identifies the correct position of the battery inside the battery compartment.
	Interference may occur in the vicinity of the device. Local regulations and precautions for other equipment in the environment should always be followed to avoid interference. The separation distance from this device to other devices complying with standard immunity requirements in IEC 60601-1-2 is minimum 0.35 m/1ft.

	Suitable for direct current only.
	In France, it is only permitted to use the device indoors.
	Consult user manual for warnings and cautions.
	Follow instructions for use.
	Consult instructions for use.
	Complies with Type BF requirements of IEC60601-1.
XXXX	Complies with Medical Devices Directive 93/42/EEC. Complies with the Radio Equipment and Telecommunications Terminal Equipment Directive 1999/5/EC.
	The installation must be carried out in accordance with Medical Electrical Systems clause 16 in IEC 60601-1 (3rd), AAMI ES60601-1 and CSA C22.2 NO. 60601-1-08-CAN/CSA. The supplementary provisions on the reliability of electro-medical systems. It is a general rule for all electrical equipment used in the proximity of the client that: • The connected equipment must comply with IEC 60601-1 (3rd).
FCC	This device complies with part 15 of the FCC rules. Operation is subject to the following two conditions: • This device must not cause harmful interference. • This device must accept any interference received, including interference that may cause undesired operation.
IC	The term"IC" before the certification/registration number signifies that the Industry Canada technical specifications were met.
	Do not reuse.
	Used in error message dialogs if software program fails. See the detailed information in the dialog box.
	Manufacturer and date of manufacture.